(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau

(43) International Publication Date
12 October 2006 (12.10.2006)

PCT

(10) International Publication Number
WO 2006/106295 A1

(51) International Patent Classification:
A61M 5/145 (2006.01)

(21) International Application Number:
PCT/GB2006/001031

(22) International Filing Date: 21 March 2006 (21.03.2006)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0507016.4 6 April 2005 (06.04.2005) GB

(71) Applicant *(for all designated States except US)*: **CILAG AG INTERNATIONAL** [CH/CH]; Landis & Gyrstrasse 1, CH-6300 (CH).

(72) Inventors; and

(75) Inventors/Applicants *(for US only)*: **BARROW-WILLIAMS, Timothy, Donald** [GB/GB]; 200 London Road, St. Albans, Hertfordshire AL1 1PL (GB). **EDINGTON, David** [GB/GB]; Apartment 3, Brisbane House, 38 Grosvenor Road, St. Albans, Hertfordshire AL1 3AE (GB).

(74) Agents: **TUNSTALL, Christopher, Stephen** et al.; Carpmaels & Ransford, 43-45 Bloomsbury Square, London WC1A 2RA (GB).

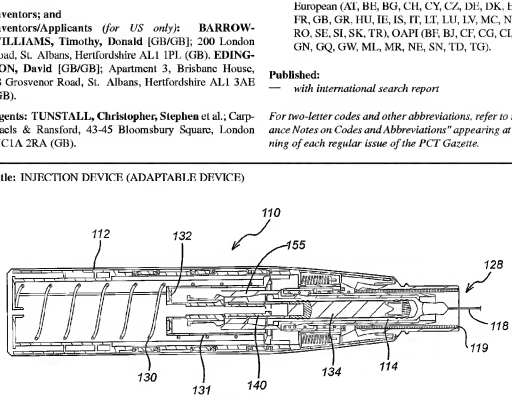
(81) Designated States *(unless otherwise indicated, for every kind of national protection available)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW.

(84) Designated States *(unless otherwise indicated, for every kind of regional protection available)*: ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:
— *with international search report*

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: INJECTION DEVICE (ADAPTABLE DEVICE)



(57) Abstract: An injection device (110) comprises a housing (112) adapted to receive a syringe (116) having a discharge nozzle (118) and a dispensing piston (114) movable in the syringe to expel the contents of the syringe through the discharge nozzle. There is also a drive (130) adapted to act on the syringe to advance it from a retracted position in which the discharge nozzle is contained within the housing to an extended position in which the discharge nozzle extends from the housing. A drive coupling (134) extends from the drive to the dispensing piston of the syringe so as to transfer movement of the drive to the piston. The drive coupling comprises a fixed-length drive coupling and an interchangeable drive coupling. There is also a method of manufacturing an injection device by assembling a first sub-assembly (210) and second sub-assembly (220) .

WO 2006/106295 A1

INJECTION DEVICE

FIELD OF THE INVENTION

The present invention relates to an injection device of the type that receives a syringe, extends it, discharges its contents and then retracts it automatically.

BACKGROUND OF THE INVENTION

Known injection devices are shown in WO 95/35126 and EP-A-0 516 473 and tend to employ a drive spring and some form of release mechanism that releases the syringe from the influence of the drive spring once its contents are supposed to have been discharged, to allow it to be retracted by a return spring.

Generally, such injection devices require a high force drive spring in order to reliably empty the syringe in the time before retraction of the syringe. When the drive spring is first released, the spring first takes up clearance in the syringe, then extends the syringe and needle and then delivers the drug. The spring force is significantly higher than that required for these initial steps and excess energy is liberated in the form of noise and vibration resulting from recoil of the spring.

An injection device can generally operate with a range of syringe fill volumes. When the fill volume is low, there can be substantial clearance to be taken up and hence louder noise and higher recoil on actuation.

It is therefore desirable to minimise the noise and recoil to avoid startling the patient for a range of syringe fill volumes.

SUMMARY OF THE INVENTION

The injection device of the present invention is designed to deal with the aforementioned problem and other issues.

In view of the foregoing and in accordance with a first aspect of the present invention, there is provided an injection device comprising:

a housing adapted to receive a syringe having a discharge nozzle and a dispensing piston movable in the syringe to expel the contents of the syringe through the discharge nozzle;

a drive adapted on activation to act on the syringe to advance it from a retracted position in which the discharge nozzle is contained within the housing to an extended position in which the discharge nozzle extends from the housing;

a drive coupling for extending from the drive to the dispensing piston of the syringe so as to transfer movement of the drive to the piston,

characterised in that the drive coupling comprises a fixed-length drive coupling and an interchangeable drive coupling.

By varying the length of the interchangeable drive coupling, the noise and vibration resulting from actuation of the device can be minimised for a given volume of conents in the syringe.

Preferably, the interchangeable drive coupling comprises a rigid member configured to connect with the dispensing piston and with the fixed-length drive coupling.

The rigid member has a longitudinal axis.

Advantageously, the quantity of the contents of the syringe which is expelled in use determines the length of the interchangeable drive coupling along its longitudinal axis.

Generally, the length of the interchangeable drive coupling is inversely proportional to the quantity of the contents of the syringe.

In one embodiment of the present invention, there is provided an interchangeable release element adapted to disengage the drive from the fixed-length drive coupling after a predetermined amount of movement of the piston.

Preferably, the interchangeable release element is a constriction adapted to act on at least one arm linking the fixed-length drive coupling to the drive, thereby releasing the drive from the fixed-length drive coupling.

Advantageously, the length of the constriction determines the predetermined amount of movement of the piston.

Generally, the length of the constriction is inversely proportional to the quantity of the contents of the syringe.

In accordance with a second aspect of the invention, there is provided a method of manufacturing an injection device, comprising:

inserting a syringe having a piston into a first sub-assembly;

inserting an interchangeable drive coupling into the syringe to contact the piston;

providing a second sub-assembly comprising a drive and a fixed-length drive coupling; and

assembling the first sub-assembly and second sub-assembly together,

wherein, on assembly, the fixed-length drive coupling and interchangeable drive coupling communicate in use to transfer movement of the drive to the piston.

Preferably, the interchangeable component has a longitudinal axis and comprises a rigid member configured to connect with the piston and with the drive coupling.

In one embodiment of the present invention, there is the additional step of selecting the interchangeable drive coupling in accordance with its length determined by the quantity of the contents of the syringe.

In a further embodiment of the present invention, there is the further step of connecting an interchangeable release element to the first sub-assembly before the step of assembling,

wherein the interchangeable release element is adapted to actuate after a predetermined amount of movement of the piston a delay mechanism in the drive acting on the fixed-length drive coupling.

Preferably, the interchangeable release element is selected in accordance with its length determined by the quantity of the contents of the syringe.

BRIEF DESCRIPTION OF THE DRAWINGS

The invention will now be described by way of example with reference to the accompanying drawings, in which:

Figure 1 shows a cross-sectional view of an injection device according to the present invention; and

Figure 2 shows an enlarged part of the injection device shown in figure 1.

Figure 3 shows an exploded view of components of the injection device according to the present invention; and

Figure 4 shows a perspective view of sub-assemblies of the injection device according to the present invention.

DETAILED DESCRIPTION OF THE DRAWINGS

Figures 1 to 4 show an injection device 110, having an injection device housing 112. The end of the housing 112 has an exit aperture 128, through which the end of a sleeve 119 can emerge.

The housing 112 contains a hypodermic syringe 114 of conventional type, including a syringe body 116 defining a reservoir and terminating at one end in a hypodermic needle

118 and at the other in a flange 120. The syringe body 116 is of substantially constant diameter along the length of the reservoir, and is of significantly smaller diameter close to the end of the syringe which terminates in the hypodermic needle. A drive coupling 134 acts through the bung of the syringe to discharge the contents of the syringe 114 through the needle 118. This drive coupling 134 constrains a drug 124 to be administered within the reservoir defined by syringe body 116. Whilst the syringe illustrated is of hypodermic type, this need not necessarily be so. Transcutaneous or ballistic dermal and subcutaneous syringes may also be used with the injection device of the present invention.

As illustrated, the syringe is housed within a syringe carrier 150. The syringe carrier 150 has a proximal end 151 through which the needle 118 of the syringe protrudes. The needle 118 is attached to the syringe body 116 of the syringe by a needle sub-assembly 172 which has a reduced diameter. At the proximal end 151 of the syringe carrier 150, there is a section of reduced diameter 173 which supports the end of the syringe 114 on its body 116. The syringe carrier 150 also includes a pair of flexible projections 152. The pair of flexible projections 152 communicate with a corresponding pair of locking apertures on a return spring support 160 so that the syringe carrier 150 cannot move relative to the return spring support 160. The syringe carrier 150 also comprises a bearing surface 153 close to its second end, against which a corresponding bearing surface of the return spring support 160 is biased by a return spring 126. The return spring 126, via the return spring support 160 and the syringe carrier 150 biases the syringe 114 from an extended position in which the needle 118 extends from the aperture 128 in the housing 112 to a retracted position in which the needle 118 is contained within the housing 112.

The syringe carrier 150 comprises a sheath 154 into which the syringe 114 can be inserted from a distal end 170. The syringe 114 is provided with a boot (not shown). If the syringe were to fail or break, the sheath 154, which surrounds the syringe 114 along its length, would contain the broken pieces of syringe and reduce the likelihood of them from escaping from the injection device 110.

The housing is further provided with a resilient latch member 161 that is biased into a position in which it engages a locking surface 163 on the return spring support 160. Before engaging the locking surface 163, the latch member 161 also extends through a latch

opening 165 in the sleeve 119. The latch member 161 includes a ramped surface 167 against which an edge of the latch opening 165 acts in the manner of a cam acting on a cam follower.

The housing also includes an actuator, and a drive which here takes the form of a compression drive spring 130. Drive from the drive spring 130 is transmitted via a multi-component drive to the piston of the syringe 114 to advance the syringe from its retracted position to its extended position and discharge its contents through the needle 118. The drive accomplishes this task by acting directly on the drug 124 and the syringe 114. Static friction between the drive coupling 134 and the syringe body 116 initially ensures that they advance together, until the return spring 126 bottoms out or the syringe body 116 meets some other obstruction (not shown) that retards its motion.

The multi-component drive between the drive spring 130 and the syringe 114 consists of three principal components. A drive sleeve 131 takes drive from the drive spring 130 and transmits it to a drive element 132. This in turn transmits drive to the drive coupling 134 already mentioned.

The drive element 132 includes a hollow stem 140, the inner cavity of which forms a collection chamber 142 in communication with a vent 144 that extends from the collection chamber through the end of the stem 140. The drive coupling 134 includes a blind bore 146 that is open at one end to receive the stem 140 and closed at the other. As can be seen, the bore 146 and the stem 140 define a fluid reservoir 148, within which a damping fluid is contained.

A trigger 214 is provided on the housing 112 remote from the exit aperture 128. The trigger, when operated, serves to decouple the drive sleeve 131 from the housing 112, allowing it to move relative to the housing 112 under the influence of the drive spring 130. The operation of the device is then as follows.

Initially, the return spring carrier 152, and consequently the syringe carrier 150 and syringe 114, are prevented from movement by the resilient latch member 161. By moving the sleeve 119 in a direction into the housing 112, the edge of the latch opening 165 is brought

into contact with the ramped surface 167 of the latch member 161, causing the latch member 161 to move outwards and thus to disengage from the return spring support 160. Once the latch member 161 has disengaged from the locking surface 163, the syringe is free to move.

The actuator is then depressed and the drive spring 130 is released. The drive spring 130 moves the drive sleeve 131, the drive sleeve 131 moves the drive element 132 and the drive element 132 moves the drive coupling 134. The drive coupling 134 moves and, by virtue of static friction and hydrostatic forces acting through the drug 124 to be administered, moves the syringe body 114 against the action of the return spring 126. The syringe body 114 moves the syringe carrier 150, which in turn moves the return spring support 160 and compresses the return spring 126. The hypodermic needle 118 emerges from the exit aperture 128 of the housing 112. This continues until the return spring 126 bottoms out or the syringe body 116 meets some other obstruction (not shown) that retards its motion. Because the static friction between the drive coupling 134 and the syringe body 116 and the hydrostatic forces acting through the drug 124 to be administered are not sufficient to resist the full drive force developed by the drive spring 130, at this point the drive coupling 134 begins to move within the syringe body 116 and the drug 124 begins to be discharged. Dynamic friction between the drive coupling 134 and the syringe body 116 and hydrostatic and hydrodynamic forces now acting through the drug 124 to be administered are, however, sufficient to retain the return spring 126 in its compressed state, so the hypodermic needle 118 remains extended.

Before the drive coupling 134 reaches the end of its travel within the syringe body 116, so before the contents of the syringe have fully discharged, flexible latch arms linking the first and drive couplings 132, 134 reach an interchangeable release element 155 connected to the distal end of the syringe carrier 150.

The interchangeable release element 155 is essentially a constriction which moves the flexible latch arms to a position so that they no longer couple the drive element 132 to the drive coupling 134. Once this happens, the drive element 132 acts no longer on the drive coupling 134, allowing the drive element 132 to move relative to the drive coupling 134.

Because the damping fluid is contained within a reservoir 148 defined between the end of the drive element 132 and the blind bore 146 in the drive coupling 134, the volume of the reservoir 146 will tend to decrease as the drive element 132 moves relative to the drive coupling 134 when the former is acted upon by the drive spring 130. As the reservoir 148 collapses, damping fluid is forced through the vent 144 into the collection chamber 142. Thus, once the flexible latch arms have been released, some of the force exerted by the drive spring 130 does work on the damping fluid, causing it to flow though the constriction formed by the vent 144; the remainder acts hydrostatically through the fluid and through friction between the first and drive couplings 132, 134, thence via the drive coupling 134. Consequently, the drive coupling 134 continues to move within the syringe body 116 and the drug 124 continues to be discharged. Losses associated with the flow of the damping fluid do not attenuate the force acting on the body of the syringe to a great extent. Thus, the return spring 126 remains compressed and the hypodermic needle remains extended.

After a time, the drive coupling 134 completes its travel within the syringe body 116 and can go no further. At this point, the contents of the syringe 114 are completely discharged and the force exerted by the drive spring 130 acts to retain the drive coupling 134 in its terminal position and to continue to cause the damping fluid to flow though the vent 144, allowing the drive element 132 to continue its movement.

Before the reservoir 148 of fluid is exhausted, flexible latch arms linking the drive sleeve 131 with the drive element 132 reach another constriction within the housing 112. The constriction moves the flexible latch arms so that they no longer couple the drive sleeve 131 to the drive element 132. Once this happens, the drive sleeve 131 acts no longer on the drive element 132, allowing them to move relative each other. At this point, the forces developed by the drive spring 130 are no longer being transmitted to the syringe 114. The only force acting on the syringe will be the return force from the return spring 126 which acts on the end of the syringe 114 nearest to the needle 118 via the return spring support 160 and the syringe carrier 150. Consequently, the syringe is returned to its retracted position and the injection cycle is complete.

As can be seen from figure 3, the drive coupling 134 comprises a fixed-length drive coupling 134a and an interchangeable drive coupling 134b. The length of the

interchangeable drive coupling 134b can be varied by changing it for a different interchangeable drive coupling. The interchangeable drive coupling 134b includes a cup 234 at its end adjacent the fixed-length drive coupling 134a for receiving a protrusion 235 on the fixed-length coupling 134b.

By varying the overall length of the drive coupling 134 by varying the length of the interchangeable drive coupling 134a, the initial start position (before actuation of the trigger) of a proximal end 240 (the end nearest the discharge needle 118 of the syringe 114) of the drive coupling 134 can be varied. Hence, the proximal end 240 can be arranged to have an initial start position as close as possible to the contents of the syringe – i.e. the initial volume of the syringe 114 can correspond substantially to the volume of the syringe contents. This way, the clearances which have to be taken up when the drive 130 is first released can be minimised, thereby reducing noise and recoil in the injection device 110.

Hence, a smaller volume of liquid in the syringe requires a longer interchangeable drive coupling 134a.

Since the length of the drive coupling 134 is variable, it follows that the point at which the flexible latch arms no longer couple the drive element 132 to the drive coupling 134 needs to be varied to ensure that the contents of the syringe can be completely discharged. This is done by varying the length of the interchangeable release element 155.

The longer the drive coupling 134 is, the longer interchangeable release element 155 needs to be.

The interchangeable release element 155 is provided with flexible arms 271 for connecting the interchangeable release element 155 to the syringe carrier 150 at cut-outs 281 on the syringe carrier 150.

As shown in figure 4, the injection device 110 can be assembled in two sub-assemblies for ease of manufacture.

A first sub-assembly 210 comprises inserting the syringe 114, syringe carrier 150, interchangeable drive coupling 134b and interchangeable release element 155.

A second sub-assembly 220 comprises the housing 112 and drive elements and actuators of the injection device 110, including the fixed length drive coupling 134a.

This way, a decision on the amount of contents to be inserted into the syringe can be made independently of the construction of the drive elements (which are difficult to assemble) because the length of interchangeable drive coupling 134b and interchangeable release element 155 can be varied immediately before final assembly of the complete injection device 110 by combining the first sub-assembly 210 with the second sub-assembly 220.

It will of course be understood that the present invention has been described above purely by way of example and modifications of detail can be made within the scope of the invention.

CLAIMS

1. An injection device comprising:

a housing adapted to receive a syringe having a discharge nozzle and a dispensing piston movable in the syringe to expel the contents of the syringe through the discharge nozzle;

a drive adapted on activation to act on the syringe to advance it from a retracted position in which the discharge nozzle is contained within the housing to an extended position in which the discharge nozzle extends from the housing;

a drive coupling for extending from the drive to the dispensing piston of the syringe so as to transfer movement of the drive to the piston,

characterised in that the drive coupling comprises a fixed-length drive coupling and an interchangeable drive coupling.

2. The injection device of claim 1, wherein the interchangeable drive coupling comprises a rigid member configured to connect with the dispensing piston and with the fixed-length drive coupling.

3. The injection device of claim 2, wherein the rigid member has a longitudinal axis.

4. The injection device of claim 3, wherein the quantity of the contents of the syringe which is expelled in use determines the length of the interchangeable drive coupling along its longitudinal axis.

5. The injection device of claim 4, wherein the length of the interchangeable drive coupling is inversely proportional to the quantity of the contents of the syringe.

6. The injection device of any one of the preceding claims, further comprising:

an interchangeable release element adapted to actuate after a predetermined amount of movement of the piston a delay mechanism in the drive acting on the fixed-length drive coupling.

7. The injection device of claim 6, wherein the interchangeable release element is a

constriction adapted to act on at least one arm linking the fixed-length drive coupling to the drive, thereby releasing the drive from the fixed-length drive coupling.

8. The injection device of claim 7, wherein the length of the constriction determines the predetermined amount of movement of the piston.

9. The injection device of claim 8, wherein the length of the constriction is inversely proportional to the quantity of the contents of the syringe.

10. A method of manufacturing an injection device, comprising:

inserting a syringe having a piston into a first sub-assembly;

inserting an interchangeable drive coupling into the syringe to contact the piston;

providing a second sub-assembly comprising a drive and a fixed-length drive coupling; and

assembling the first sub-assembly and second sub-assembly together,

wherein, on assembly, the fixed-length drive coupling and interchangeable drive coupling communicate in use to transfer movement of the drive to the piston.

11. The method of claim 10, wherein the interchangeable component has a longitudinal axis and comprises a rigid member configured to connect with the piston and with the drive coupling.

12. The method of claim 10 or claim 11, further comprising selecting the interchangeable drive coupling in accordance with its length determined by the quantity of the contents of the syringe.

13. The method of any one of claims 10 to 12, further comprising connecting an interchangeable release element to the first sub-assembly before the step of assembling, wherein the interchangeable release element is adapted to actuate after a predetermined amount of movement of the piston a delay mechanism in the drive acting on the fixed-length drive coupling.

14. The method of any one of claims 10 to 13, further comprising selecting the

interchangeable release element in accordance with its length determined by the quantity of the contents of the syringe.

15. The injection device of any one of the preceding claims, substantially as hereinbefore described with reference to the accompanying drawings.

16. The method of any one of the preceding claims, substantially as hereinbefore described with reference to the accompanying drawings.

FIG. 1

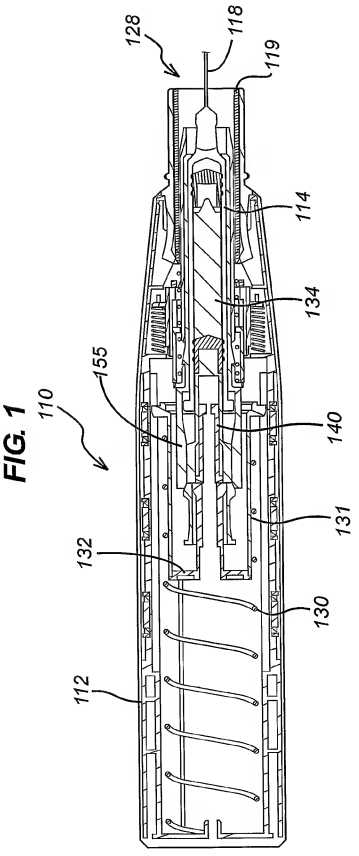
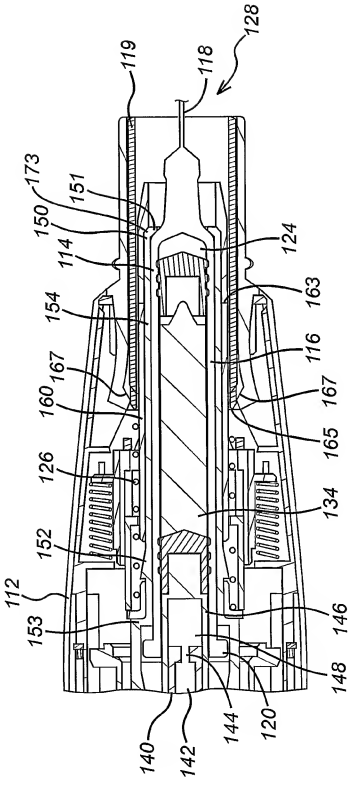


FIG. 2


128
119
118
173
151
150
114
124
154
163
116
167
160
167
165
126
134
152
112
153
146
148
140
142
144
120

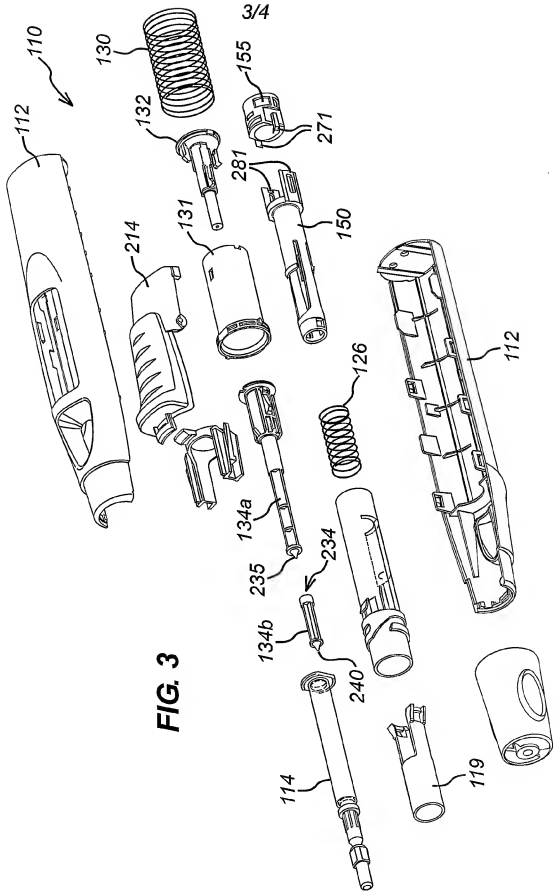

110
130
155
112
132
271
281
214
131
150
126
112
134a
235
234
134b
240
114
119

FIG. 3

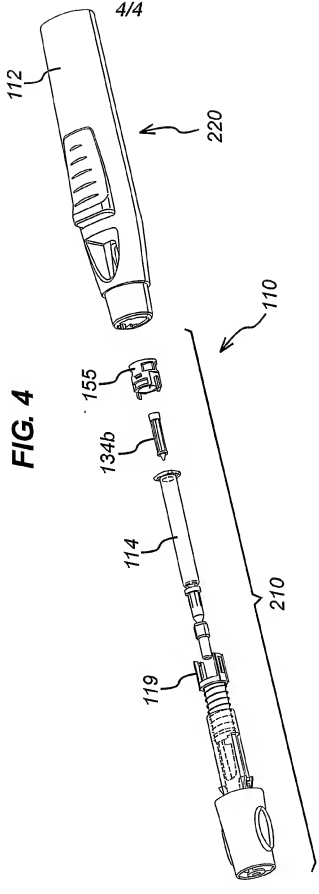


FIG. 4

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2006/001031

A. CLASSIFICATION OF SUBJECT MATTER
INV. A61M5/145

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
A61M

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 02/056947 A (MEDRAD, INC; DEDIG, JAMES, ALBERT; LAZZARO, FRANK, A; COWAN, KEVIN, P;) 25 July 2002 (2002-07-25) abstract; figure 7	1-5, 10-13
X	US 5 520 653 A (REILLY ET AL) 28 May 1996 (1996-05-28) abstract; figures	1,10
A	EP 0 482 677 A (MALLINCKRODT MEDICAL, INC; MALLINCKRODT, INC) 29 April 1992 (1992-04-29) abstract; figures	1,10
A	EP 1 518 575 A (MATSUSHITA ELECTRIC INDUSTRIAL CO., LTD) 30 March 2005 (2005-03-30) abstract; figures 2,3	1,10

-/--

[X] Further documents are listed in the continuation of Box C. [X] See patent family annex.

* Special categories of cited documents :

"A" document defining the general state of the art which is not considered to be of particular relevance
"E" earlier document but published on or after the international filing date
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
"O" document referring to an oral disclosure, use, exhibition or other means
"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
18 May 2006	30/05/2006

Name and mailing address of the ISA/	Authorized officer
European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016	Ehrsam, F

Form PCT/ISA/210 (second sheet) (April 2005)

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2006/001031

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	GB 2 414 402 A (* CILAG AG INTERNATIONAL) 30 November 2005 (2005-11-30) abstract; figures	1,10

Form PCT/ISA/210 (continuation of second sheet) (April 2005)

INTERNATIONAL SEARCH REPORT

International application No.
PCT/GB2006/001031

Box II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☒ Claims Nos.: 15,16
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210

3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.

2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.

3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box II.2

Claims Nos.: 15,16

Since claims 15 and 16 are referring to the figures of he present application they are unclear since no meaningful search could be executed.

The applicant's attention is drawn to the fact that claims relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the claims are amended following receipt of the search report or during any Chapter II procedure. If the application proceeds into the regional phase before the EPO, the applicant is reminded that a search may be carried out during examination before the EPO (see EPO Guideline C-VI, 8.5), should the problems which led to the Article 17(2) declaration be overcome.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
PCT/GB2006/001031

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 02056947	A	25-07-2002	CN	1494444 A	05-05-2004
			EP	1351730 A1	15-10-2003
			JP	2005504557 T	17-02-2005
US 5520653	A	28-05-1996	AT	225674 T	15-10-2002
			AT	304377 T	15-09-2005
			DE	69624251 D1	14-11-2002
			DE	69624251 T2	18-06-2003
			DE	69635194 D1	20-10-2005
			EP	0847287 A1	17-06-1998
			JP	11512016 T	19-10-1999
			JP	3471812 B2	02-12-2003
			WO	9709077 A1	13-03-1997
EP 0482677	A	29-04-1992	AT	80045 T	15-09-1992
EP 1518575	A	30-03-2005	CN	1665557 A	07-09-2005
			WO	2004004809 A1	15-01-2004
			US	2005209569 A1	22-09-2005
GB 2414402	A	30-11-2005	WO	2005115509 A1	08-12-2005